Atty Dkt. No.: INTM-035 USSN: (unassigned)

I. AMENDMENTS

AMENDMENTS TO THE CLAIMS

Cancel claims 3, 4, 7-12, and 18-44 without prejudice to renewal.

Please enter the amendments to claims 1, 5, 6, and 14-17, as shown below.

Please enter new claims 45-50, as shown below.

- 1. (Currently amended) A method of treating a patient suffering from idiopathic pulmonary fibrosis, comprising administering to the patient an effective amount of IFN-γ, where the patient has a forced vital capacity (FVC) that is at least about 55% of the predicted normal value and/or has a carbon monoxide diffusing capacity (DL_{CO}) that is at least about 30% [[35%]] of the predicted normal value.
- 2. (Original) A method of increasing the probability of survival of a patient suffering from idiopathic pulmonary fibrosis (IPF), comprising administering to the patient an effective amount of IFN- γ , where the patient has a forced vital capacity (FVC) that is at least about 55% of the predicted normal value and/or has a carbon monoxide diffusing capacity (DL_{CO}) that is at least about 35% of the predicted normal value.

3.-4. (Canceled)

- 5. (Currently amended) The method of claim 1, wherein the patient has a DL_{CO} that is at least 35% A method of treating a patient suffering from idiopathic pulmonary fibrosis, comprising administering to the patient an effective amount of IFN γ, where the patient has a forced vital capacity (FVC) that is at least about 55% of the predicted normal value and/or has a carbon monoxide diffusing capacity (DL_{CO}) that is at least about 30% of the predicted normal value.
- 6. (Currently amended) The method of <u>claim 1</u> any one of claims 1-5, wherein IFN-γ is administered to the patient for a period [[of]] <u>selected from</u> about 48 weeks, about 60 weeks, about one year, about 70 weeks, about 93 weeks, about 2 years, and the remainder of the patient's life.

7.-12, (Canceled)

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13. (Currently amended) The method of <u>claim 1</u> any one of claims 1-5, wherein the method further comprises administering a corticosteroid to the individual.

- 14. (Currently amended) The method of claim 1 any one of claims 1-5, wherein IFN- γ is administered in a dose of from about 1.48 µg/kg body weight to about 4.4 µg/kg body weight about 80 µg/m² to about 90 µg/m².
- 15. (Currently amended) The method of claim 1 any one of claims 1-5, wherein IFN- γ is administered in a dose of about 200 µg.
- 16. (Currently amended) The method of <u>claim 1</u> any one of claims 1-5, wherein IFN-γ is administered three times weekly.
- 17. (Currently amended) The method of <u>claim 1</u> any one of claims 1-5, wherein IFN-γ is administered by subcutaneous administration.

18.-44. (Canceled)

- 45. (New) The method of claim 2, wherein IFN-γ is administered to the patient for a period selected from about 48 weeks, about 60 weeks, about one year, about 70 weeks, about 93 weeks, about 2 years, and the remainder of the patient's life.
- 46. (New) The method of claim 2, wherein the method further comprises administering a corticosteroid to the individual.
- 47. (New) The method of claim 2, wherein IFN- γ is administered in a dose of from about 1.48 µg/kg body weight to about 4.4 µg/kg body weight.
 - 48. (New) The method of claim 2, wherein IFN-γ is administered in a dose of about 200 μg.

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49. (New) The method of claim 2, wherein IFN-γ is administered three times weekly.

50. (New) The method of claim 2, wherein IFN- γ is administered by subcutaneous administration.